



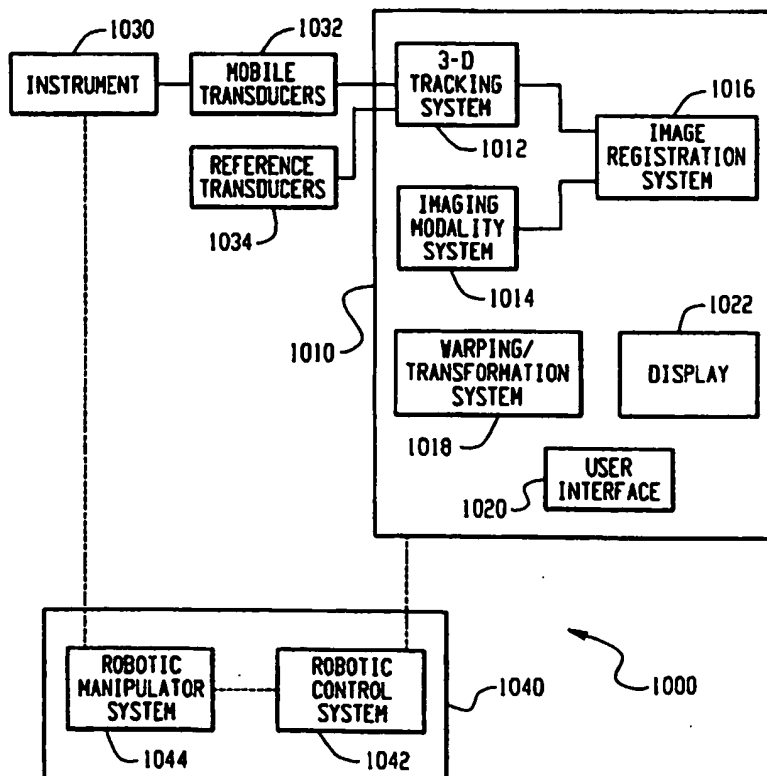
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(54) Title: **SYSTEM FOR CARRYING OUT SURGERY, BIOPSY AND ABLATION OF A TUMOR OR OTHER PHYSICAL ANOMALY**

(57) Abstract

A system for carrying out surgery on a bodily structure (e.g., breast and liver) with greater precision, accuracy, comfort and minimal invasiveness. A 3-dimensional tracking and imaging system (1000) is used to obtain an accurate position of an instrument as it is maneuvered by an operator, and to mark a location on the subject bodily structure. The system is particularly advantageous for surgical procedures for biopsying and destroying tumors in bodily structures which are easily deformable.



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**SYSTEM FOR CARRYING OUT SURGERY,
BIOPSY AND ABLATION OF A TUMOR OR
OTHER PHYSICAL ANOMALY**

Related Applications

The present application is a continuation-in-part (CIP) of co-pending International Application No. PCT/CA96/00194, filed March 26, 1996, which is a continuation-in-part (CIP) of U.S. Application Serial No. 08/411,959, filed March 28, 1995, now U.S. Patent No. 5,515,853. The contents of both applications are incorporated herein by reference.

Field of Invention

The present invention relates generally to a system for carrying out a surgical procedure on a bodily structure (e.g., breast, liver, pancreas, kidney, uterus or other solid organ), and more particularly to a system for tracking an instrument within a bodily structure, marking the location of a tumor, and biopsying or destroying the tumor.

Background of the Invention

Breast biopsies are currently performed using a device known as a Core Biopsy System. The Core Biopsy system first obtains a stereo-mammogram from a patient's breast, while the breast is immobilized by being compressed between two plates, and uses these two images to calculate the 3-D coordinates of the suspected

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tumor. A needle is then fired into the breast and a biopsy is taken of the suspected tumor. If the biopsy is positive, then the patient is scheduled for tumor removal surgery. It should be noted that before the biopsy procedure is commenced, the tumor needs to be manually identified by a radiologist.

5 The surgical procedure generally proceeds in the following manner. A patient undergoes multi-plane mammography, a radiologist examines the film, and then inserts a wire into the breast so that it punctures the tumor. This procedure is visualized using repetitive x-ray imaging. More recently, the stereotactic breast imaging system has been used to localize the tumor more precisely and assist in the
10 insertion of the wire. The patient is then sent to the operating room, and the breast is prepared for surgery by the application of a topical sterilant. The surgeon then cuts the breast open, following the wire until the lesion is found and excised.

 One of the undesirable factors of the foregoing procedure is the presence of a long wire through the breast for many hours at a time while awaiting surgery. This is
15 highly traumatic for the patient and undesirable. Secondly, during surgery, the surgeon must follow the wire into the breast. Since this may not be the optimal trajectory, the surgeon would ideally like to plan the entry pathway independent of the wire, or eliminate the wire altogether. This can be done only if the location of the lesion within the breast can be identified using a system that takes into account the
20 inherent deformability of the breast tissue. It should be appreciated that the problem associated with the deformability of breast tissue applies equally to other easily deformable bodily structures such as the liver.

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The current problem limiting use of stereotactic breast surgery is the large difference between the position and shape of the breast during mammography and surgery. In this regard, images taken during mammography become unusable for stereotactic positioning during the surgical procedure. While stereotactic surgery can be done with the breast compressed, and the patient lying on the stereotactic table, this is not desirable. The ideal way to do this surgery is with the patient on her back, as is done routinely.

The present invention overcomes these and other drawbacks of prior art systems and provides a system having significantly improved accuracy and providing greater comfort to the patient.

Summary of the Invention

According to the present invention there is provided a system for performing stereotactic surgery with improved accuracy and comfort.

It is an object of the present invention to provide a system for performing surgery on a bodily structure in a minimally invasive manner.

It is another object of the present invention to provide a system for performing surgery on a bodily structure which is inherently deformable.

It is another object of the present invention to provide a system for performing surgery, wherein there is increased precision of the localization of a tumor, enabling more precise surgery.

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It is still another object of the present invention to provide a system for performing surgery, wherein a constant 3-D reference frame is established.

It is still another object of the present invention to provide a system for performing surgery, wherein deformation of the subject bodily structure does not alter a 3-D frame of reference.

It is yet another object of the present invention to provide a system for performing breast surgery in which there is greatly reduced potential for disfiguring or altering the shape of the breast.

It is yet another object of the present invention to provide a system for performing surgery of the liver, or other internal organ, in which there is more accurate locating of a tumor.

It is yet another object of the present invention to provide a system for performing surgery of the liver, or other organ, in which there is a need for more accurate locating of a surgical instrument.

Still other objects and advantages of the invention will become apparent to those skilled in the art upon a reading and understanding of the following detailed description, accompanying drawings and appended claims.

Brief Description of the Drawings

The invention may take physical form in certain parts and arrangements of parts, a preferred embodiment and method of which will be described in detail in this

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specification and illustrated in the accompanying drawings which form a part hereof, and wherein:

Fig. 1 shows a 3-dimensional tracking and imaging system according to a preferred embodiment of the present invention.

5 Fig. 2 shows a breast having a reference transducer deposited therein while the breast is under compression;

Fig. 3 shows a transducer arrangement according to a preferred embodiment of the present invention;

10 Figs. 4A and 4B illustrates the progression of a probe through a breast shown against the original stereo mammograms;

Fig. 5A shows reference transducers and a "homing beacon" transducer located on a breast;

Fig. 5B shows the scene of Fig. 5A, as shown in 3-D on a display unit;

15 Fig. 6A shows a probe having a transducer in contact with the breast shown in Fig. 4A;

Fig. 6B shows the scene of Fig. 6A, as shown in 3-D on a display unit;

Fig. 7 shows a liver having a plurality of reference transducers applied thereto;

Fig. 8 illustrates a procedure for generating a 3-D scene including a 2-D ultrasound image of the liver shown in Fig. 7.

20 Fig. 9 shows insertion of a locatable tube into the liver shown in Fig. 7; and

Fig. 10 shows the insertion of a cryoprobe into a trackable sleeve located at the site of a tumor.

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Detailed Description of the Preferred Embodiment

It is known that using the time-of-flight principle of high frequency sound waves, it is possible to accurately measure distances within an aqueous medium, such as inside the body of a living being during a surgical procedure. High frequency sound, or ultrasound, is defined as vibrational energy that ranges in frequency from 100 kHz to 10 MHz. The device used to obtain three-dimensional measurements using sound waves is known as a sonomicrometer. Typically, a sonomicrometer consists of a pair of piezoelectric transducers (i.e., one transducer acts as a transmitter while the other transducer acts as a receiver). The transducers are implanted into a medium, and connected to electronic circuitry. To measure the distance between the transducers, the transmitter is electrically energized to produce ultrasound. The resulting sound wave then propagates through the medium until it is detected by the receiver.

The transmitter typically takes the form of a piezoelectric crystal that is energized by a high voltage spike, or impulse function lasting under a microsecond. This causes the piezoelectric crystal to oscillate at its own characteristic resonant frequency. The envelope of the transmitter signal decays rapidly with time, usually producing a train of six or more cycles that propagate away from the transmitter through the aqueous medium. The sound energy also attenuates with every interface that it encounters.

The receiver also typically takes the form of a piezoelectric crystal (with similar characteristics to the transmitter piezoelectric crystal) that detects the sound

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energy produced by the transmitter and begins to vibrate in response thereto. This vibration produces an electronic signal in the order of millivolts, that can be amplified by appropriate receiver circuitry.

The propagation velocity of ultrasound in an aqueous medium is well documented. The distance traveled by a pulse of ultrasound can therefore be measured simply by recording the time delay between the instant the sound is transmitted and when it is received. Three-dimensional coordinates can be determined from the distance measurement.

Referring now to the drawings wherein the showings are for the purposes of illustrating a preferred embodiment of the invention only and not for purposes of limiting same, Fig. 1 shows a three-dimensional (3-D) tracking and imaging system 1000 for use in connection with the procedure of the present invention. 3-D tracking and imaging system 1000 is generally comprised of a computer system 1010, mobile transducers 1032, reference transducers 1034, an instrument 1030 and an optional robotics subsystem 1040.

Computer system 1010 is generally comprised of a 3-D tracking system 1012, an imaging modality system 1014, an image registration system 1016, a warping and geometry transformation system 1018 ("warp system"), a user interface 1020 and a display 1022. It should be appreciated that 3-D tracking system 1012 may take the form of a sound-based system or an electromagnetic-based system. Both time of flight and phase relationships may be used to determine distance. Preferably, 3-D tracking system 1012 takes the form of the 3-D ultrasound tracking system described

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in U.S. Patent No. 5,515,853 and PCT Application No. WO96/31753, both of which are incorporated herein by reference.

Instrument 1030 may take the form of a catheter, a probe (e.g., a cryoprobe), a sensor, a needle, a scalpel, a forceps or other device or instrument used in a surgical or diagnostic procedure. Mobile transducers 1032 and reference transducers 1034 may take the form of an ultrasonic transducer or an electronic transducer. However, for purpose of illustrating a preferred embodiment of the present invention, transducers 1032 and 1034 will take the form of ultrasonic transducers (i.e., piezoelectric crystals).

A plurality of mobile transducers 1032 are fitted to instrument 1030. One or more reference transducers 1034 provide a reference position relative to mobile transducers 1032. In this respect, reference transducers 1034 may be located to provide an internal reference frame inside a patient's body or on the surface of a patient body to provide an external reference frame.

As indicated above, reference transducers 1034 may be transmitters, transceivers or receivers that can generate ultrasound or electromagnetic radiation, that can be detected by mobile transducers 1032.

3-D tracking system 1012 transforms the multiple distance measurements between all of the transducers 1032, 1034 into XYZ coordinates relative to a referenced axis, as described in detail above. It should be appreciated that the reference frame provided by reference transducers 1034 must be self-determining, that is, if the reference frame becomes distorted, this distortion needs to be detected by

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reference transducers 1034. Detection is typically done by using transceivers that can determine the distance between any combination of two transducers, and hence their relative spacial coordinates in 3-D space. In this regard, the position of the transducers is obtained in 3-D from the images acquired of the bodily structure (e.g., tissue or organ) that show "dots" where the transducers are located, and also from the transducers themselves when they are in the bodily structure. If there is some discrepancy in the distances between all combinations of transducers, then the bodily structure must have deformed (i.e., "warped") after the images were acquired. A mathematical coordinate transformation can be used to specify exactly how to correct the image set and account for the warping. The distance between any combination of two transducers is determined by having each transducer send a signal to all other transducers. In this way, all the distances between the transducers are known. From these distances, XYZ coordinates can be calculated, in reference to some transducer as the origin.

Imaging modality system 1014 acquires 2-D, 3-D or 4-D image data sets from an imaging source, such as fluoroscopy, an MRI (magnetic resonance imaging), CT (computerized tomography) or 2-D or 3-D ultrasound device, to provide a "template" through or against which the shape, position and movement of instrument 1030 being tracked can be displayed. The template typically takes the form of an image of the environment surrounding the instrument (e.g., a bodily structure). It should be noted that if multiple (3-D) volumes are acquired at different time intervals, a 4-D image is obtained (i.e., 3-D image changing over time).

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Image registration system 1016 registers the position of instrument 1030 within the spatial coordinates of the image data set provided by imaging modality system 1014. The position of instrument 1030 is provided by the 3-D tracking system 1012. Image registration system 1016 will provide a display of instrument 1030 at its proper 3-D location inside the bodily structure and orientation relative to the bodily structure itself. It should be appreciated that registration system 1016 may be user assisted, or completely automated if image processing algorithms are implemented to automatically detect the spacial locations of the transducers (typically the reference transducers) in the image data set.

Warp system 1018 is a software-based system that transforms or "warps" the image data sets by the appropriate values to correspond to a deformation that has occurred in the reference frame between the time that the image data set were acquired and the time that the procedure is to be implemented during surgery. Accordingly, warp system 1018 is typically comprised of a matrix transformation routine that maps the deformed geometry onto the original image data set, and distorts it appropriately.

User interface 1020 enables a user to interact with computer system 1010, including programming computer system 1010 to perform a desired function. For example, a particular view for display can be selected. Instruments 1030 (e.g., probes or catheters) can be activated using user interface 1020. Display 1022 displays to the user registered images provided by image registration system 1016.

Optional robotics system 1040 is generally comprised of a robotics control system 1042 and a robotic manipulator system 1044. Robotics control system 1042

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controls robotic manipulator system 1044 to follow a programmed path that can be appropriately changed, based on shifting, warping or changes in the shape of a bodily structure at the time of surgery. Robotic manipulator system 1044 physically moves instrument 1030 as instructed by robotic control system 1042.

5 The above-described 3-D tracking and imaging system 1000 can be used to provide both stereotactic localization during biopsy and surgery that is more interactive than existing stereotaxy table systems, and tagging of a tumor so that it can subsequently be localized during conventional surgery. It should be appreciated that while the present invention is described with reference to the biopsy and ablation of a
10 tumor, it is also suitable for use in connection with other physical anomalies.

 Tagging of a tumor located in a breast will now be described with reference to Fig. 2. A plurality of external reference transducers 20 are affixed to the surface of breast 10. Reference transducers 20 provide a stereotactic external reference frame for the interactive 3-D display of the movement of probe 40 during insertion of internal
15 transducer 30, as will be described below. Tumor 12 is tagged by inserting internal ultrasonic transducer 30 into tumor 12 during conventional mammography, wherein breast 10 is placed under compression by the use of compression plates 8. Transducer 30 takes the place of the localizer needle that is presently inserted into the tumor according to prior art methods.

20 Reference transducers 20 may take the form of individual stick-on elements, or part of an adhesive strip. Fig. 3 shows an exemplary embodiment of an arrangement for affixing reference transducer 20 using an adhesive. Reference transducer 20 is

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supported by an adhesive patch 24. A matching gel 26 is applied to the adhesive patch 24 and reference transducer 20 is arranged therein. Gel 26 provides acoustic coupling. Electrical leads 22 of reference transducer 20 exit through an opening in adhesive patch 24.

5 It should be appreciated that reference transducers 20 will appear on the two mammograms obtained from at two slightly different angles, and can be used to generate fiducial markers for the stereographic determination of 3-D coordinates of tumor 12, relative to these markers. Moreover, the motion of probe 40 can also be referenced against these bi- plane mammograms using transducer 30. Accordingly, a
10 user can track the motion of probe 40 both in a 3-D viewing environment, as well as against the original radiograms, during the deposition of transducer 30, which will act as a "homing beacon" for the tumor during subsequent biopsy or surgery. The progression of an instrument through a breast is shown against mammograms in Figs. 4A and 4B.

15 Once transducer 30 has been deposited in tumor 12, and probe 40 removed from the breast 10, the patient can comfortably walk around, since electrical leads 32 connected to transducer 30 are very flexible, and can be taped to the patient's skin. It should be appreciated that transducer 30 can reliably denote the location of tumor 12 during subsequent surgery since it remains lodged in tumor 12 by means of small
20 barbs that deploy during insertion.

 The foregoing approach is a significant improvement over conventional stereotaxy. Since a tremendous change occurs in the shape of the breast between

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mammography and surgery, the tumor may be in a completely different location than previously perceived during mammography. As a result, any stereotactic registration of external breast shape with the internal breast images is lost. However, since the tumor is tagged with transducer 30, its location can always be determined during subsequent procedures.

After the foregoing procedure, the patient goes to the operating room, the breast is prepared for surgery, and new adhesive, with ultrasonic transducers imbedded therein, are attached to the skin. Transducers 20 and 30 are connected to the 3-D tracking and imaging system described above. It should be appreciated that transducers 20 and 30 enable the tracking of additional transducers that may be inserted into the breast during subsequent surgery.

Fig. 5A shows breast 10 having tumor 12 tagged with internal transducer 30. The attachment of external reference transducers 20 and the presence of internal transducer 30 in tumor 12 enables the generation of a 3-D viewing environment within which tumor 12 can be localized relative to breast 10 (Fig. 5B). This means of visualizing the spatial location of tumor 12 relative to the outside of breast 10 is important in planning the tumor removal surgery. The conventional technique is to simply follow a previously inserted wire into the breast. This is not desirable since it may not be the most cosmetically desirable path to take. By analyzing a 3-D display, such as shown in Fig 5B, a surgeon can decide from which direction to begin incisions.

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As the incisions are made, a secondary probe 50 can be touched into the wound to determine if the trajectory should be modified. Fig 6A shows probe 50 touching breast 10. It should be noted that probe 50 has a transducer 52 arranged on its tip. Accordingly, a 3-D display as shown in Fig. 6B can be generated. Once
5 transducer 52 mounted to probe 50 comes into contact with the tissue, it will appear in the 3-D display and can be localized relative to transducer 30 (i.e., "homing beacon"). Accordingly, the 3-D display allows the surgical path to be visualized and corrected, as necessary. Because the external reference frame formed by reference transducers 20 is affixed to the external surface of breast 10, it does not matter if the
10 breast tissue deforms following mammogram imaging. In this regard, transducer 30 will always be shown relative to the new configuration of transducers 20 affixed to the external surface of breast 10. Moreover, since external reference transducers 20 communicate with each other, they will set up a new, changing coordinate frame regardless to what extent the breast tissue is manipulated. In each case, the relative
15 position of transducer 30 is displayed within this coordinate system.

The foregoing approach provides numerous advantages and overcomes the current limitations of stereotactic surgery that is based on a fixed coordinate system. With regard to clinical advantages, breast surgery can be done much less invasively. When developed further, it may be possible to remove tumors in a much less invasive
20 way, by inserting small catheters that ablate, suction or in some other way destroy the tumor without having to open up the breast for visual inspection. This will greatly reduce the potential for disfiguring or altering the shape of the breast. Another

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important clinical advantage is the increased precision of the localization of the tumor, enabling more precise surgery. The theoretical precision of the 3-D tracking system described above is 20 μ m. In practice, the 3-D spatial precision is less than 1 mm, and due to geometrical transformation errors, this may rise to 2 mm. This is apparently still better than the precision currently available using the wire insertion technique or conventional stereotactic tables. Such increased precision will improve the success rate of tumor excision, without the need for larger, more radical lumpectomies. Yet another clinical advantage is that the foregoing procedure can be readily applied to both biopsies, open surgery or fully closed, minimally invasive surgery. It provides a real-time, user assisted means of approaching tumors inside the breast and is much less costly than conventional stereotactic equipment.

The present invention may also be used to locate an instrument at a tumor site, according to another embodiment of the present invention. Fig. 7 shows a liver 60 having a tumor 14. A 3-dimensional reference frame of liver 60 is established by attaching a plurality of reference transducers 90 to the external surface of liver 60. In this regard, transducers 90 may be attached under laparoscopic guidance. Next, the location of tumor 14 is determined using guided 2-dimensional ultrasound 110, which generates a 2-D ultrasound imaging plane 112. A 2-D ultrasound imaging plane is displayed within the 3-D reference frame to form a 3-D scene. When tumor 14 is transected by ultrasound imaging plane 112, the user places a cursor mark in the 3-D scene to identify the center of tumor 14. The 3-D tracking and imaging system determines the 3-D coordinates of tumor 14 within the 3-D scene, relative to the 3-D

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reference frame established by transducers 90. It should be appreciated that since transducers 90 are fixed to liver 60, the location of tumor 14 remains fixed relative to the 3-D reference frame, even as liver 60 itself is manipulated causing deformation of liver 60.

5 Referring now to Figs. 9 and 10 a cryoprobe 100 (or other instrument) is located at tumor 14. In this respect, an ultrasonically locatable sheath attachment 70 is inserted into liver 60 and positioned such that its end corresponds to the location of tumor 14. Sheath attachment 70 preferably takes the form of a hollow rigid sleeve or tube having transducers 80 mounted thereto. A preferred sheath attachment 70 is
10 described in detail in co-pending U.S. Application Serial No. _____, entitled "Tracking Data Sheath", filed March 7, 1997, and incorporated herein by reference. Transducers 80 allow the position of sheath attachment 70 to be tracked using the 3-D tracking and imaging system, described above.

Once sheath attachment 70 has been located at the site of tumor 14, cryoprobe
15 100 is inserted into sheath attachment 70, such that it stops at the mouth of the sheath. Accordingly, cryoprobe 100 does not require physical modifications to be located at the site of tumor 14. Next, sheath attachment 70 is pulled back along the shaft of cryoprobe 100. Thereafter, cryoprobe 100 is energized to ablate tumor 14 in a well known manner.

20 Alternatively, the foregoing procedure can be performed by inserting a trackable blunt guide instrument having ultrasonic transducers mounted thereto and locating the guide instrument at the site of the tumor. Next, a tube or sleeve is placed

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at the site of the tumor by placing the tube over the guide instrument. Then, the guide instrument is removed, leaving just the sleeve. A cryoprobe (or other instrument) is then inserted through the sleeve, thus locating the cryoprobe at the site of the tumor.

The invention has been described with reference to a preferred embodiment.

5 Obviously, modifications and alterations will occur to others upon a reading and understanding of this specification. It is intended that all such modifications and alterations be included insofar as they come within the scope of the appended claims or the equivalents thereof.

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Having thus described the invention, it is now claimed:

1. A method for carrying out surgery on a tumor in a bodily structure for the purpose of biopsy or ablation, comprising:

mounting a plurality of reference transducer means to fixed positions relative to the bodily structure to provide a reference frame;

5 obtaining a template image of the bodily structure;

registering the reference frame with the template image to obtain a 3-dimensional scene;

locating a beacon transducer means at the location of the tumor by displaying the position of the beacon transducer means within the 3-dimensional scene, the

10 beacon transducer means identifying the location of the tumor; and

locating a first instrument means at the location of the tumor by displaying the position of the first instrument means relative to the reference frame and beacon transducer means.

2. A system according to claim 1, wherein said step of mounting a plurality of reference transducer means to fixed positions includes the step of mounting a plurality of transducer means to the surface of a breast.

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3. A system according to claim 1, wherein the step of mounting a plurality of reference transducer means to a fixed positions includes the step of mounting a plurality of transducer means to the external surface of a solid organ.

4. A system according to claim 1, wherein said template image is a mammogram.

5. A system according to claim 1, wherein said template image is a radiological image.

6. A system according to claim 1, wherein the step of locating the beacon transducer means at the location of the tumor includes the steps of:

attaching the beacon transducer means to a second instrument means;

moving the second instrument means to the location of the tumor; and

5 detaching the beacon transducer means from the second instrument means at the location of the tumor.

7. A system for carrying out surgery on a tumor in a bodily structure for the purpose of biopsy or ablation, comprising:

a plurality of reference transducer means located at fixed positions on the bodily structure to provide a reference frame;

5 means for obtaining a template image of the bodily structure;

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registration means for registering the reference frame with the template image to obtain a 3-dimensional scene;

display means for displaying the 3-dimensional scene of the bodily structure;

10 first instrument means for locating a beacon transducer means at the site of the tumor, the beacon transducer means identifying the location of the tumor; and

second instrument means for performing a surgical procedure at the site of the tumor, wherein the second instrument means has a transducer means for indicating the position of the first instrument means relative to the reference frame and the location of the beacon transducer means.

8. A system according to claim 7, wherein said fixed positions are on the surface of a breast.

9. A system according to claim 7, wherein said fixed positions are on the external surface of a solid organ.

10. A system according to claim 7, wherein said template image is a mammogram.

11. A system according to claim 7, wherein said template image is a radiological image.

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12. A system according to claim 7, wherein said first instrument means includes means for separating said beacon transducer means from said first instrument means.

13. A system according to claim 7, wherein each of said plurality of reference transducer means comprises:

a transducer; and

support means for supporting the transducer, said support mean including an adhesive for affixing the reference transducer means to the fixed position on the
5 bodily structure and a gel for acoustic coupling.

14. A system according to claim 13, wherein said transducer is a piezoelectric crystal transducer.

15. A method for carrying out surgery on a tumor in a bodily structure for the purpose of biopsy or ablation, comprising:

mounting a plurality of reference transducer means to fixed positions on the bodily structure to provide a reference frame;

5 obtaining a template image of the bodily structure;

registering the reference frame with the template image to obtain a 3-dimensional scene;

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locating a trackable sleeve means at the location of the tumor by displaying the position of the sleeve means within the 3-dimensional scene; and

10 locating a first instrument means at the location of the tumor by positioning the first instrument means within the sleeve means.

16. A method according to claim 15, wherein the step of mounting a plurality of transducer means to fixed positions includes the step of mounting a plurality of transducer means to a solid organ under laparoscopic guidance.

17. A method according to claim 15, wherein said template image is a 2-dimensional ultrasound image.

18. A method according to claim 15, wherein said method further comprises the step of displaying a reference mark in the 3-dimensional scene, at the location of the tumor to facilitate locating the trackable sleeve means at the location of the tumor.

19. A method according to claim 15, wherein the method further includes the step of removing the trackable sleeve from the first instrument means.

20. A system for carrying out surgery on an anomaly in a bodily structure for the purpose of biopsy or ablation, comprising:

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a plurality of reference transducer means mounted at fixed positions on the
bodily structure to provide a reference frame;

5 means for obtaining a template image of the bodily structure;

registration means for registering the reference frame with the template image
to obtain a 3-dimensional scene;

trackable sleeve means locatable relative to an associated anomaly to identify
a location of the associated anomaly;

10 display means adapted for displaying the position of the trackable sleeve

means within the 3-dimensional scene to locate the sleeve means at the location of the
associated anomaly; and

first instrument means adapted for performing a surgical procedure at the
location of the associated anomaly, the first instrument means located at the location
15 of the associated anomaly by positioning the first instrument means within the sleeve
means.

21. A system according to claim 20, wherein said fixed position is the
external surface of a solid organ.

22. A system according to claim 20, wherein said template image is a 2-
dimensional ultrasound image.

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23. A system according to claim 20, wherein said system further comprised means for displaying a reference mark in the 3-dimensional scene at the location of the anomaly to facilitate locating the trackable sleeve means at the location of the anomaly.

24. A system according to claim 20, wherein said first instrument means is a cryoprobe.

25. A method for carrying out surgery on a tumor in a bodily structure for the purpose of biopsy or ablation, comprising:

mounting a plurality of reference transducer means to fixed positions to provide a reference frame;

5 obtaining a template image of the bodily structure;

registering the reference frame with the template image to obtain a 3-dimensional scene;

locating a trackable instrument means at the location of the tumor by displaying the position of the trackable sleeve means within the 3-dimensional scene;

10 arranging a sleeve means over the trackable instrument means;

removing the trackable instrument means from the location of the tumor; and

inserting a second instrument means into the sleeve means.

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26. A system according to claim 25, wherein said second instrument means is an ablation means.

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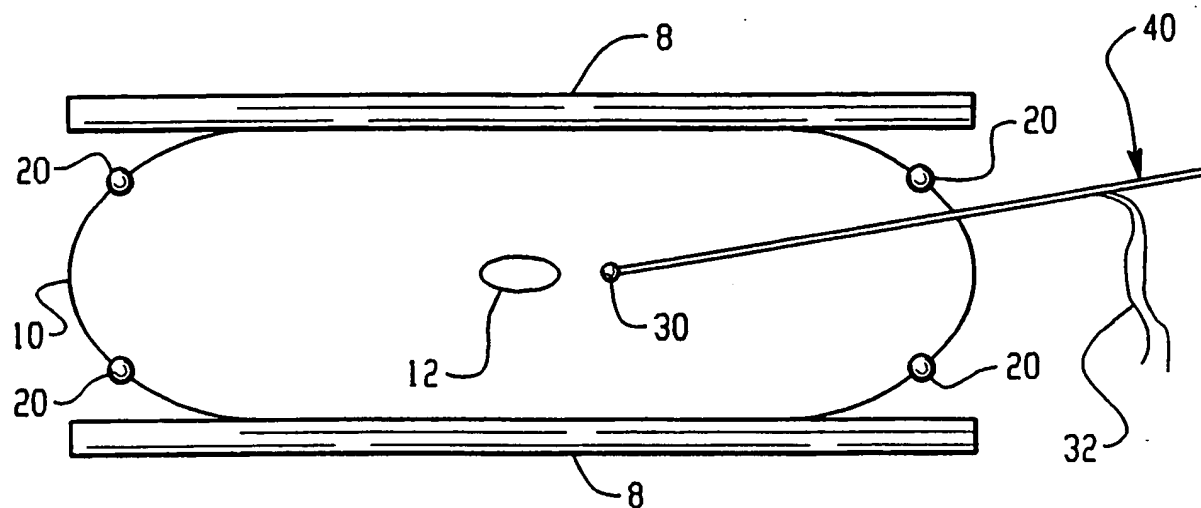


Fig. 2

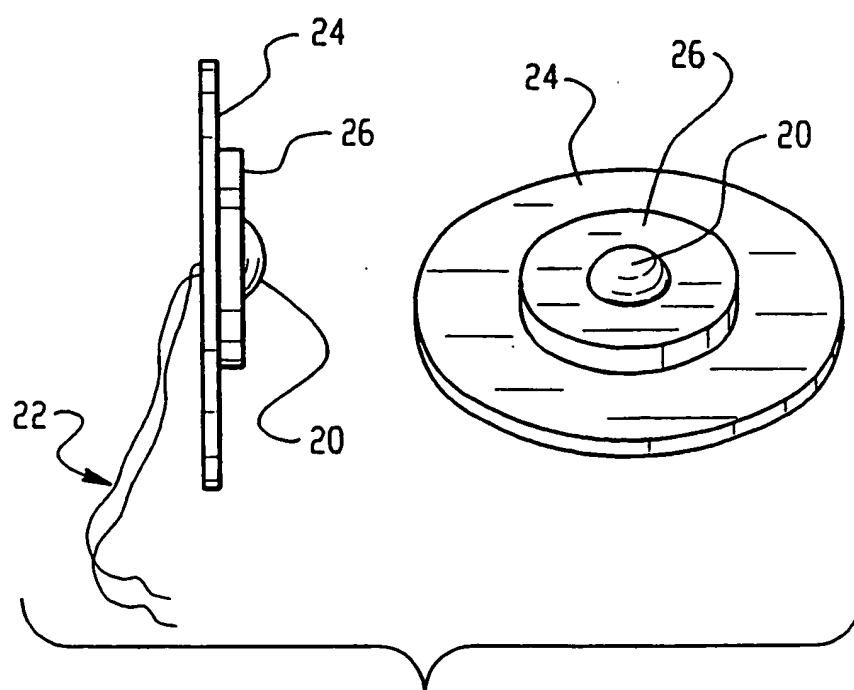


Fig. 3

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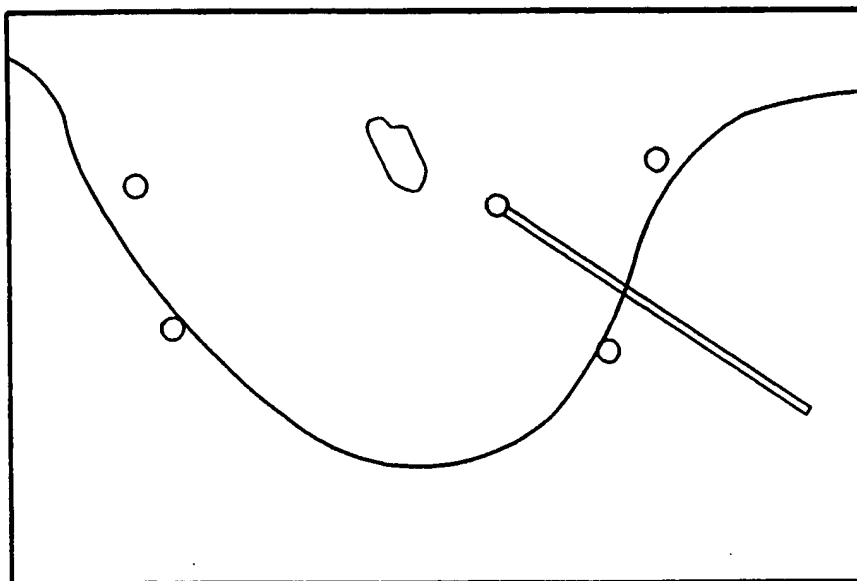


Fig.4A

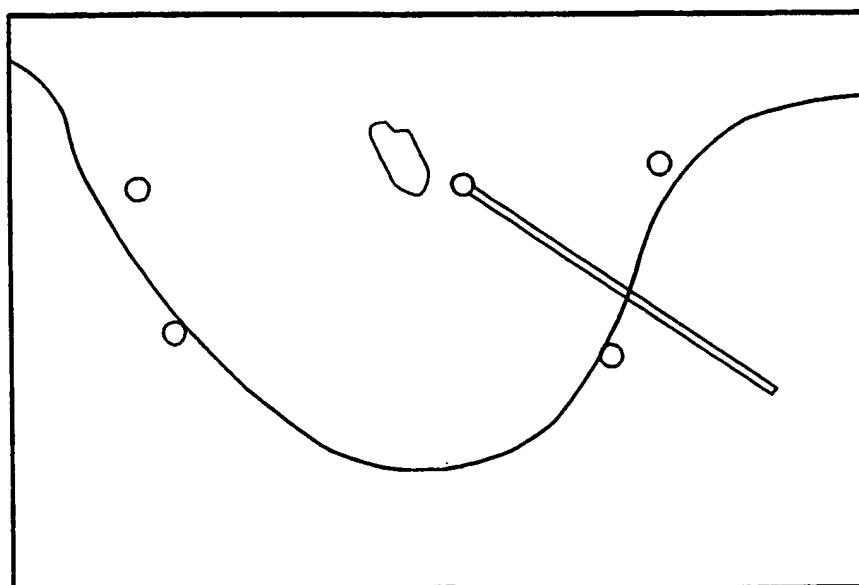
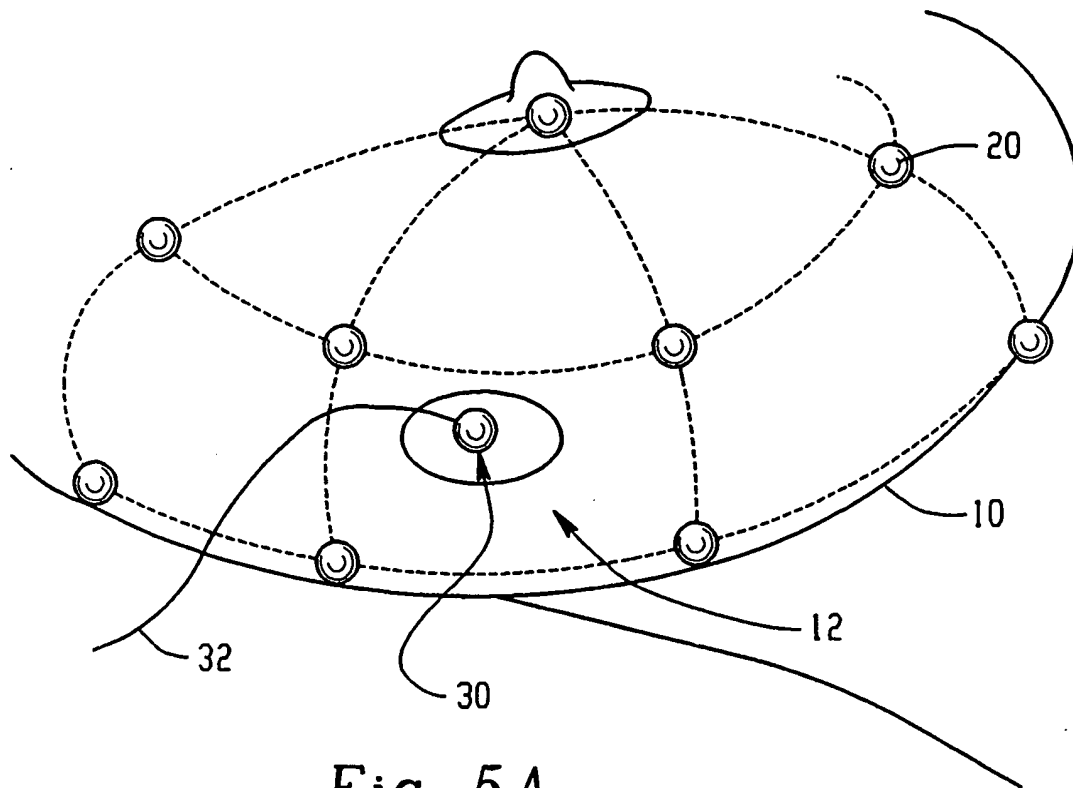
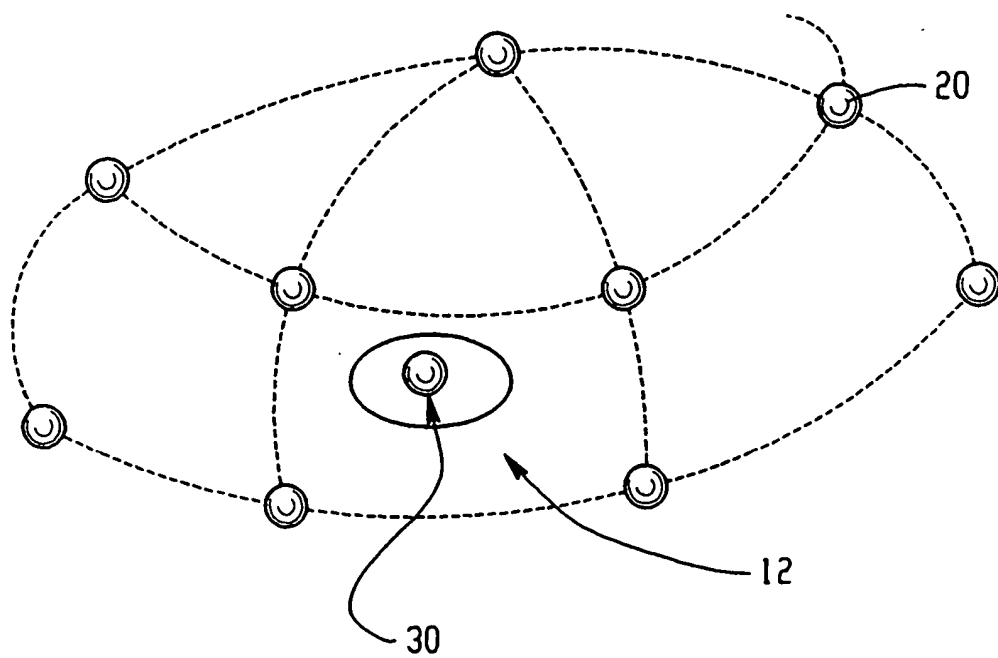


Fig.4B

*Fig. 5A*

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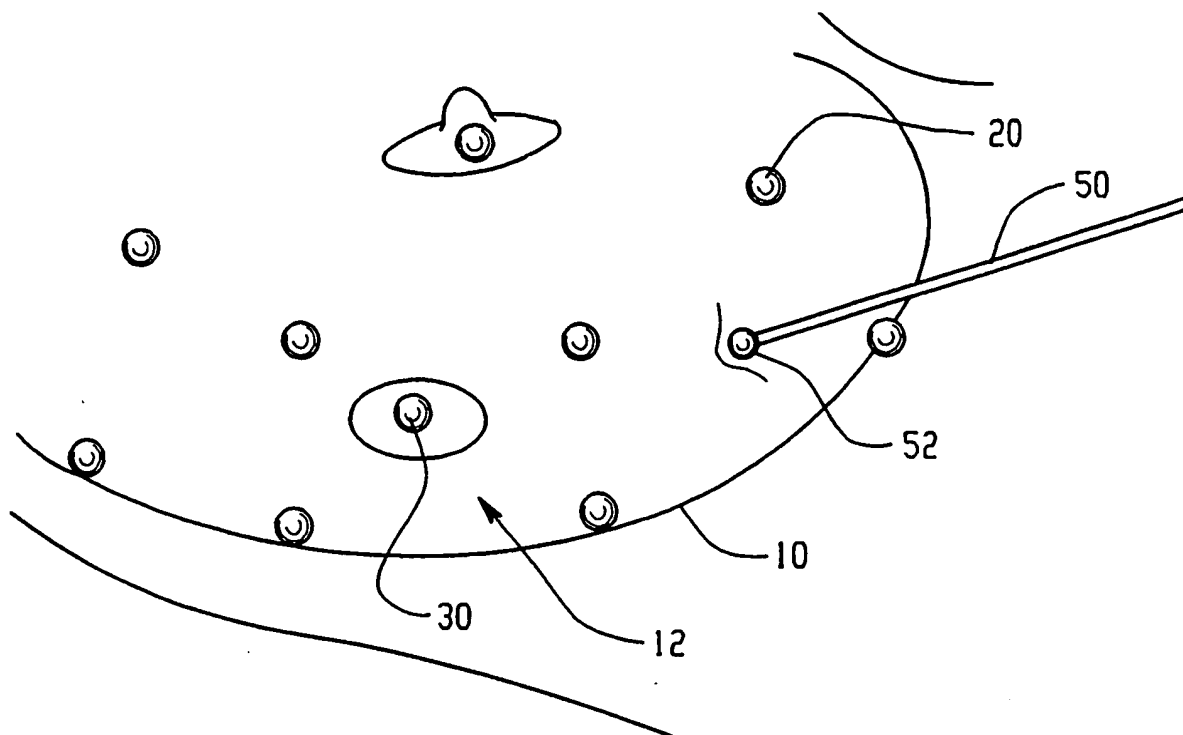


Fig. 6A

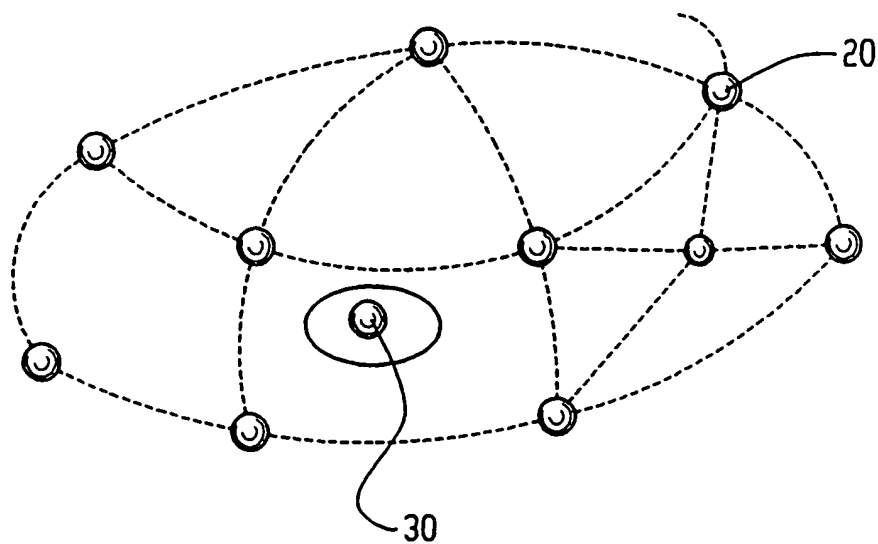


Fig. 6B

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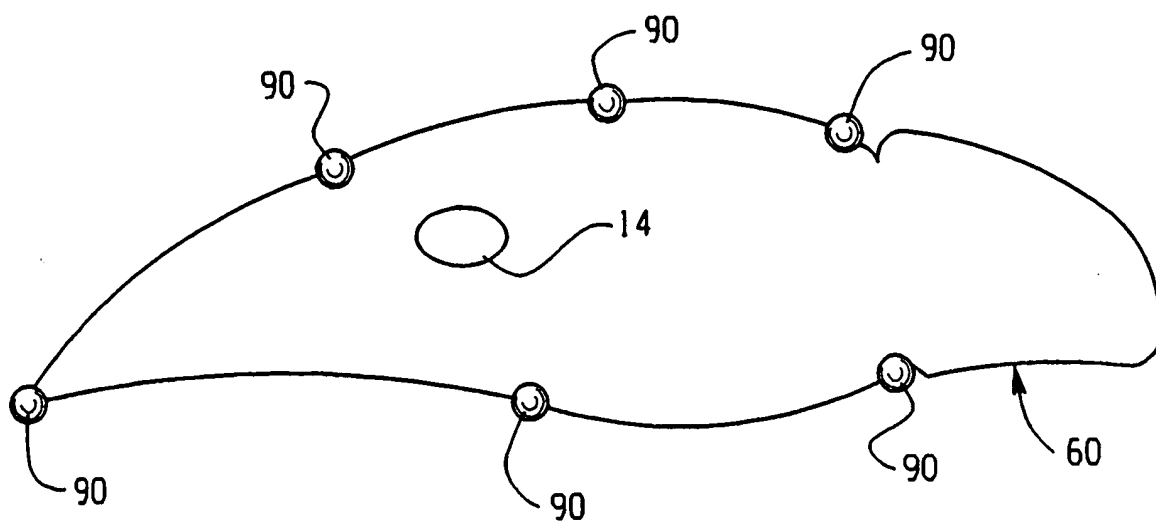


Fig. 7

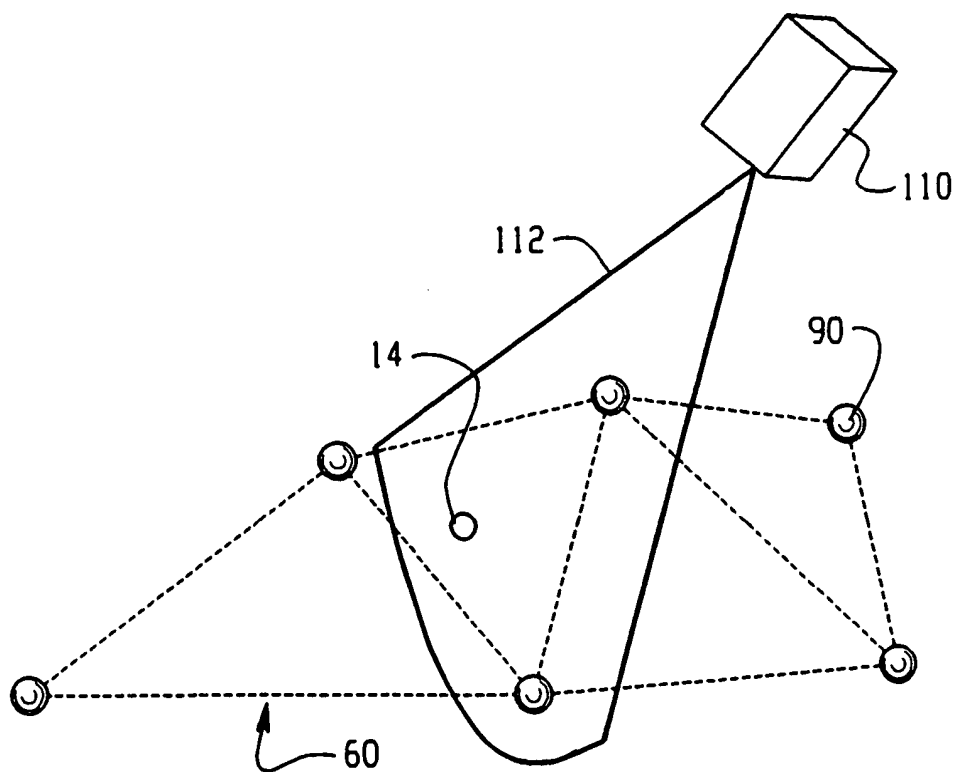


Fig. 8

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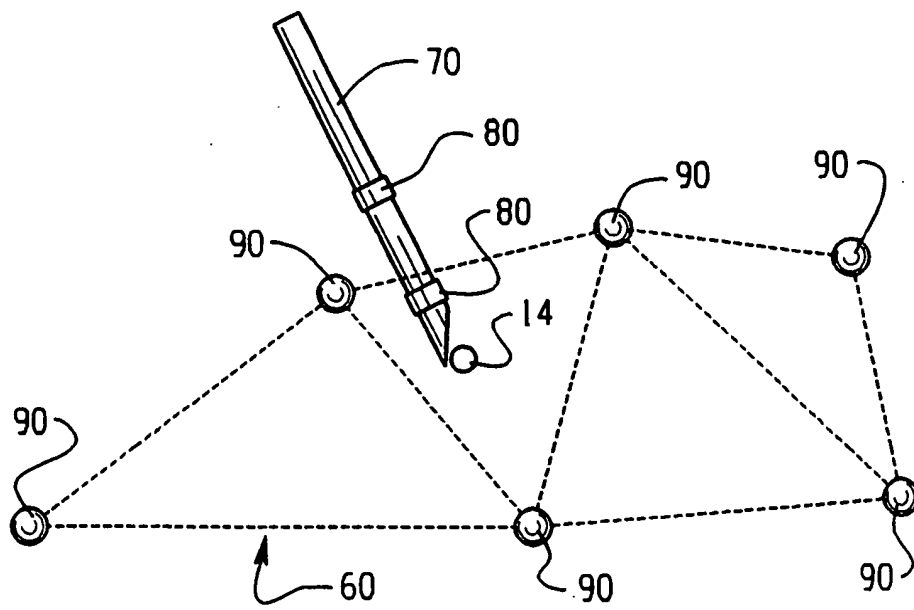


Fig. 9

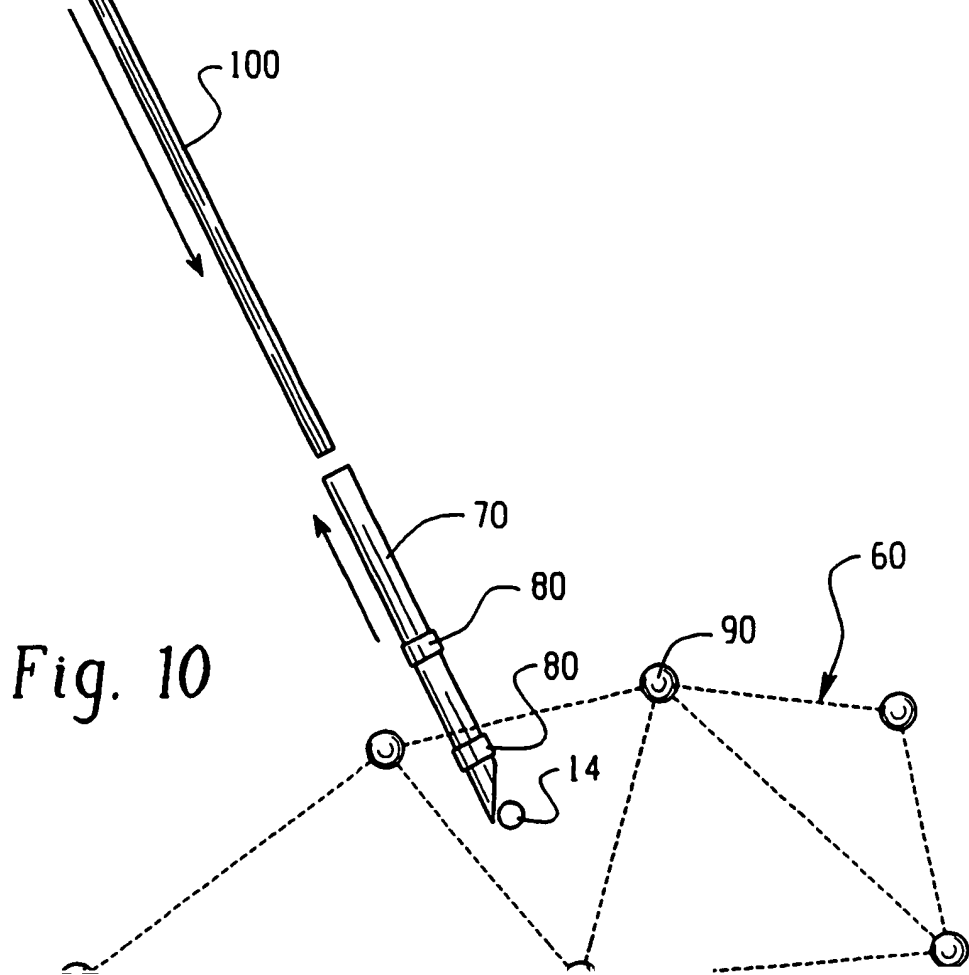


Fig. 10

INTERNATIONAL SEARCH REPORT

Intern. Application No

PCT/IB 98/00731

A. CLASSIFICATION OF SUBJECT MATTER

IPC 6 A61B19/00

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 6 A61B

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

| Category * | Citation of document, with indication, where appropriate, of the relevant passages | Relevant to claim No. |
|------------|--|-----------------------|
| Y | US 5 515 853 A (SONOMETRICS) 14 May 1996 cited in the application see column 15, line 12-52; claim 1 --- | 2-14, 20-24,26 |
| Y | EP 0 728 446 A (CLEVELAND CLINIC FOUNDATION) 28 August 1996 see column 4, line 21 - column 5, line 31 --- | 2-14, 20-24,26 |
| A | WO 96 08208 A (BIOPSY MEDICAL, INC.) 21 March 1996 see page 23, line 15-17; claim 1 --- | 2,4,6,7, 10,20,23 |
| A | US 5 260 985 A (MOSBY) 9 November 1993 see column 6, line 23-61 --- | 2,4,7, 10,20 |
| | -/-- | |



Further documents are listed in the continuation of box C.



Patent family members are listed in annex.

* Special categories of cited documents :

A document defining the general state of the art which is not considered to be of particular relevance

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O document referring to an oral disclosure, use, exhibition or other means

P document published prior to the international filing date but later than the priority date claimed

T later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

X document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

Y document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

Z document member of the same patent family

Date of the actual completion of the international search

23 July 1998

Date of mailing of the international search report

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
NL - 2280 HV Rijswijk

Authorized officer

INTERNATIONAL SEARCH REPORT

International Application No

PCT/IB 98/00731

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

| Category * | Citation of document, with indication, where appropriate, of the relevant passages | Relevant to claim No. |
|------------|--|-----------------------|
| A | WO 96 00037 A (ZUPKAS) 4 January 1996 see claims 1,10 --- | 2,7,20, 24 |
| A | US 4 706 681 A (BREYER ET AL.) 17 November 1987 see claim 1 --- | 2,7,20 |
| A | US 5 309 913 A (KORMOS ET AL.) 10 May 1994 see column 3, line 7 - column 4, line 5 ----- | 2,7,20 |

INTERNATIONAL SEARCH REPORT

International application No.
PCT/IB 98/00731

Box I Observations where certain claims were found unsearchable (Continuation of Item 1 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☒ Claims Nos.: 1,15-19,25
because they relate to subject matter not required to be searched by this Authority, namely:
Rule 39.1(iv) PCT - Method for treatment of the human or animal body by surgery
2. ☐ Claims Nos.:
because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:
3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of Item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
- ☐ No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/IB 98/00731

| Patent document cited in search report | Publication date | Patent family member(s) | Publication date |
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